

Opis choroby *

Definicja

A rare glomerular disease, histologically characterized by thickening of the capillary wall, with immune deposits predominantly containing IgG4 and C3 on the sub-epithelial side, and typically manifesting with nephrotic syndrome.

Dane

Klasyfikacja

Choroba

Synonimy

Idiopathic membranous glomerulonephritis

Primary membranous nephropathy

Idiopathic membranous glomerulonephritis

Primary membranous nephropathy

Kod ORPHA

97560

Kod OMIM

614692

Kod ICD10

N04.2

Kod ICD11

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*Źródło

orphanet